

## Adiabatic work:

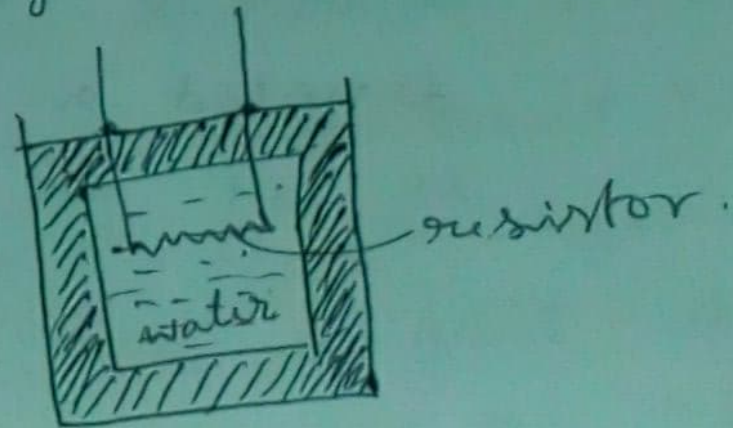
Let us consider a closed system completely surrounded by an adiabatic wall and still the system is coupled to the surroundings so that work may be done. Consider a

composite electrical system composed of a resistor immersed in water. The initial state is characterized by the

thermodynamic coordinates

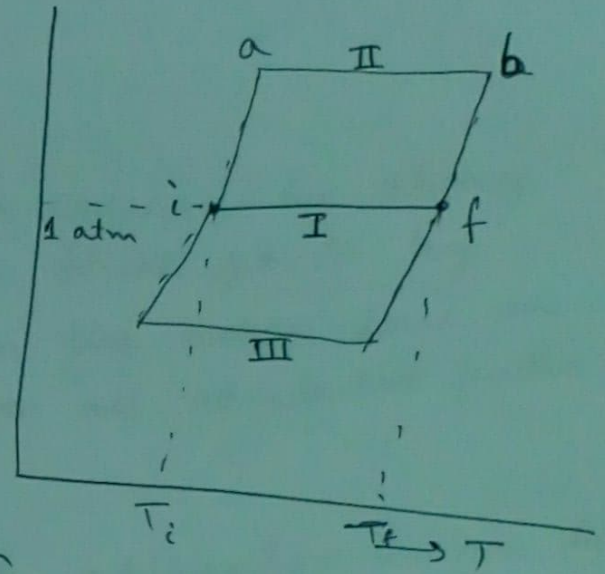
$T_i = 287.7 \text{ K} (14.5^\circ\text{C})$  and the

initial pressure  $p_i = 1 \text{ atm}$  and the final state characteri



zed by the co-ordinates  $P_f = 1 \text{ atm}$  and  $T_f = 288.7 \text{ K}$   
(15.5°C)

To take the system from  $i$  to  $f$  ~~thru~~ by performing ~~via~~ adiabatic work only through path I, it is necessary



to pass current for certain interval of time though the water when it is maintained at one atmospheric pressure.

But path I is not the only way to take the system from state  $i$  to  $f$  using adiabatic work. The same change can take place via path II also where system is first compressed from  $i$  to  $a$ , ~~a~~ then current is ~~pass~~ passed and then  $b$  to  $f$  it is again ~~compressed~~ expanded. Or, we might use a similar adiabatic path III. There are an infinite number of paths by which a system may be transferred from an initial state to a final state by the performance of adiabatic

work only. The adiabatic work is the same along all such paths, i.e. adiabatic work is path independent.

So, if a closed system is caused to change from an initial state to final state by adiabatic means only, then the work done on the system is the same for all adiabatic paths connecting the two states.

This path independence of adiabatic work helps to draw an important conclusion that there must exist an energy function which will depend only on the thermodynamic coordinates of the system like the gravitational potential energy in mechanics (this is function of the position coordinates of the system) or the electrical potential energy in ~~electro~~ electricity. This energy function is called internal-energy function and usually denoted by  $U$ . So, we have

$$W_{i \rightarrow f} \text{ (adiabatic)} = U_f - U_i, \text{ where the signs}$$

are such that if positive work is done on the system,  $U_f$  will be greater than  $U_i$ . This internal energy function is a function of thermodynamic

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variables. The physical interpretation of the difference  $U_f - U_i$  is the increase in internal energy of the system. It states that there exists an energy function whose difference between two ~~set~~ values is the energy change of the system.

For a closed hydrostatic system, the equilibrium states can be completely determined by ~~any~~ <sup>any</sup> ~~any~~ <sup>any</sup>  $p, V, T$ . The two of the three thermodynamic variables. The third is fixed by the equation of state. ~~The~~  $p, V, T$ . The internal energy function  $U$  can be regarded as a function of any two only out of the three variables. One should ~~remember~~ remember that it may not always be possible to write internal energy function in simple mathematical form, especially when one deals with real systems.

If two states of a system differs infinitesimally, then the infinitesimal change in internal energy is denoted by  $du$ .  $du$  is an exact differential since it is the differential of a state function. So, the integral of  $du$  is independent of the path between the initial and final states. In the case of hydrostatic system, if  $U$  is regard

as a function of  $T$  and  $V$ , then

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$$U = U(T, V)$$

$$\Rightarrow dU = \left( \frac{\partial U}{\partial T} \right)_V dT + \left( \frac{\partial U}{\partial V} \right)_T dV$$

or regarding  $U$  as a different function of  $T$  and  $p$ , i.e.,

$$U = U(T, p)$$

$$dU = \left( \frac{\partial U}{\partial T} \right)_p dT + \left( \frac{\partial U}{\partial p} \right)_T dp$$

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